

BSgenome.Hsapiens.UCSC.hg38

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Full genome sequences for Homo sapiens (UCSC version hg38)

Description

Full genome sequences for Homo sapiens (Human) as provided by UCSC (hg38, Dec. 2013) and stored in Biostrings objects.

Note

This BSgenome data package was made from the following source data files:

hg38.2bit from <http://hgdownload.cse.ucsc.edu/goldenPath/hg38/bigZips/>

See [?BSgenomeForge](#) and the BSgenomeForge vignette (`vignette("BSgenomeForge")`) in the **BSgenome** software package for how to make a BSgenome data package.

Author(s)

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See Also

- [BSgenome](#) objects and the [available.genomes](#) function in the **BSgenome** software package.
- [DNAString](#) objects in the **Biostrings** package.
- The BSgenomeForge vignette (`vignette("BSgenomeForge")`) in the **BSgenome** software package for how to make a BSgenome data package.

Examples

```
BSgenome.Hsapiens.UCSC.hg38
genome <- BSgenome.Hsapiens.UCSC.hg38
seqlengths(genome)
genome$chr1 # same as genome[["chr1"]]
```

```
## -----
## Extract the upstream sequences
```

```
## -----
## The upstream sequences located in
## http://hgdownload.cse.ucsc.edu/goldenPath/hg38/bigZips/
## are based on RefSeq genes (RefSeq Genes track in the Genome Browser).
## Upstream sequences based on UCSC genes (UCSC Genes track in the
## Genome Browser) can easily be extracted from the full genome
## sequences with:

library(TxDb.Hsapiens.UCSC.hg38.knownGene)
knownGene_txdb <- TxDb.Hsapiens.UCSC.hg38.knownGene
knownGene_up1000seqs <- extractUpstreamSeqs(genome, knownGene_txdb)

## Or, to get upstream sequences based on RefSeq genes:

refGene_txdb <- makeTxDbFromUCSC("hg38", "refGene")
refGene_up1000seqs <- extractUpstreamSeqs(genome, refGene_txdb)

## Note that you can make a TxDb object from various annotation
## resources. See the makeTxDbFromUCSC(), makeTxDbFromBiomart(), and
## makeTxDbFromGFF() functions in the GenomicFeatures package for more
## information.
## IMPORTANT: Make sure you use a TxDb package (or TxDb object) that
## contains a gene model based on hg38 or on a compatible genome (i.e.
## a genome with sequences identical to the sequences in hg38). See
## ?extractUpstreamSeqs in the GenomicFeatures package for more
## information.

## -----
## Genome-wide motif searching
## -----
## See the GenomeSearching vignette in the BSgenome software
## package for some examples of genome-wide motif searching using
## Biostrings and the BSgenome data packages:
if (interactive())
  vignette("GenomeSearching", package="BSgenome")
```

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